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(54) Title: WHITE MINERAL OIL-BASED DRILLING FLUID

(57) Abstract

A wellbore fluid (e.g., a drilling, completion, packer, or fracturing fluid) comprises (a) at least one additive selected from the group consisting of emulsifiers, wetting agents, viscosifiers, weighting agents, fluid loss control agents, proppants for use in hydraulically fracturing subterranean formations, and particulate agents for use in forming a gravel pack; and (b) a non-toxic white mineral oil having (i) an API gravity at 15.6 °C (60 °F) greater than 35, (ii) a content of compounds containing 14 or more carbon atoms of at least about 95 weight percent, and (iii) a pour point of at least about -30 °C (-22 °F).

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WO 94/28086

. WHITE MINERAL OIL-BASED DRILLING FLUID BACKGROUND

The present invention relates to oil-based wellbore fluids (e.g., oil-based drilling fluids) and 5 systems and processes for using them in a subterranean formation in oil and gas recovery operations.

Oil-based drilling fluids and techniques for drilling boreholes in subterranean formations to recover hydrocarbons (e.g., oil and gas) are well known to those 10 skilled in the art. In addition, a number of less toxic oil-based drilling fluids have been proposed as evidenced by, for example, U.S. Patent 4,787,990, U.S. Patent 5,096,883, U.S. Patent 5,189,012, International Publication WO 83/02949, and IADC/SPE 11355. While a number of 15 commercially available, low toxicity base oils are inexpensive, they have not acquired FDA approval for direct use in food. On the other hand, the dimer of 1-decene has obtained such FDA approval, but its prohibitive expense significantly restricts its use in drilling operations.

20 SUMMARY OF THE INVENTION

According to one aspect of the invention, there is provided a wellbore fluid comprising at least one additive selected from the group consisting of emulsifiers, wetting agents, viscosifiers, weighting agents, fluid loss 25 control agents, proppants for use in hydraulically fracturing subterranean formations, and particulate agents for use in forming a gravel pack; and a white mineral oil having (i) an API gravity at 15.6°C (60°F) greater than 35, (ii) a content of compounds containing 14 or more carbon 30 atoms of at least about 95 weight percent, and (iii) a pour point of at least about -30°C (-22°F).

According to another aspect of the invention, there is provided a drilling system comprising at least one subterranean formation; a borehole penetrating a portion of 35 at least one of the subterranean formations; a drill bit suspended in the borehole; and a drilling fluid located in the borehole and proximate the drill bit, wherein the drilling fluid is the wellbore fluid of claim 1 and at least one additive is selected from the group consisting of

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emulsifiers, wetting agents, viscosifiers, weighting agents, and fluid loss control agents.

According to further aspect of the invention, there is provided a method for drilling a borehole in a 5 subterranean formation, the method comprising the steps of: rotating a drill bit at the bottom of the borehole; and introducing a drilling fluid into the borehole (a) to pick up drill cuttings and (β) to carry at least a portion of the drilling cuttings out of the borehole, wherein the 10 drilling fluid is the wellbore fluid of claim 1 and at least one additive is selected from the group consisting of emulsifiers, wetting agents, viscosifiers, weighting agents, and fluid loss control agents.

Accordingly, there is a need for an oil-based 15 drilling fluid that employs an inexpensive, readily available, FDA approved, non-toxic base oil. The present invention satisfies this need by providing a drilling fluid comprising (a) at least one drilling fluid additive (e.g., an emulsifier, a viscosifier, a weighting agent, and an 20 oil-wetting agent) and (b) a white mineral oil having (i) an API gravity at 15.6°C (60°F) greater than 35, (ii) a content of compounds containing 14 or more carbon atoms of at least about 95 weight percent, and (iii) a pour point of at least about -30°C (-22°F).

In addition, a drilling system and a method for drilling a borehole are also provided by the invention. The drilling system comprises (a) at least one subterranean formation, (b) a borehole penetrating a portion of at least one of the subterranean formations, (c) a drill bit 30 suspended in the borehole, and (d) the above drilling fluid located in the borehole and proximate the drill bit.

Regarding the method for drilling a borehole of the present invention, this method comprises the steps of (a) rotating a drill bit at the bottom of the borehole and 35 (b) introducing the aforesaid drilling fluid into the borehole (i) to pick up drill cuttings and (ii) to carry at least a portion of the drill cuttings out of the borehole.

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DETAILED DESCRIPTION OF THE INVENTION

The white mineral oil employed as the base oil in the drilling fluid of the present invention commonly has a characterization factor greater than 12, more commonly at 5 least about 12.1, even more commonly at least about 12.2, and most commonly at least about 12.3. Generally, the characterization factor of the white mineral oil is less than about 13, more typically less than about 12.9, even more typically less than about 12.8, and most typically 10 less than about 12.7. (As used in the specification and claims, the characterization factor is determined in accordance with the procedure set forth in Watson et al., Journal of Industrial and Engineering Chemistry, 27(12):1460 et seq. (December, 1935; this publication being 15 incorporated herein in its entirety by reference), with the average boiling point being the 50 percent temperature in the Engler distillation.)

Regarding API gravity, the white mineral oil often has an API gravity at about 15.6°C (60°F) (as 20 measured by ASTM D 1298) of at least about 35.25, more often at least about 35.5, even more often at least about 35.75, and most often at least about 36. Frequently, the white mineral oil has an API gravity at about 15.6°C (60°F) of less than about 50, more frequently less than about 45, even more frequently less than about 44.75, and most frequently less than about 44.5.

The n-paraffinic content of the white mineral oil (as determined by the procedure set forth in Attachment A) is usually greater than about 5.25, more typically greater than about 5.5, even more commonly greater than about 5.75, and most frequently greater than about 6, weight percent. In some white mineral oils, the n-paraffinic content is even greater than about 10, about 15, or even about 20 weight percent.

The white mineral oil commonly has a total paraffinic content (as determined by UTM 407-90) of at least about 25, more commonly at least about 30, even more commonly at least about 33, and most commonly at least about 35, weight percent. Typically, the total paraffinic

content of the white mineral oil is less than about 70, more typically less than about 65, even more typically less than about 60, and most typically less than about 58, weight percent.

Concerning naphthenic content, the mononaphthenic content of the white mineral oil (as determined by UTM 407-90) is frequently about 20 to about 50, more frequently about 25 to about 45, even more frequently about 27 to about 44, and most frequently about 29 to about 43, weight percent. The white mineral oil often has a polynaphthenic content (as determined by UTM 407-90) of about 5 to about 30, more often about 10 to about 25, even more often about 12 to about 23, and most often about 13 to about 22, weight percent. The total naphthenic content of the white mineral oil (as determined by UTM 407-90) is typically about 30 to about 75, more typically about 35 to about 70, even more typically about 40 to about 67, and most typically about 42 to about 65, weight percent.

with respect to the carbon content of the
compounds contained in the white mineral oil, the
concentration of compounds containing 14 or more carbon
atoms (as determined by ASTM D 2887) is generally at least
about 95, more typically at least about 96, even more
commonly at least about 97, and most frequently at least
25 about 98, weight percent. Compounds comprising 18 or more
carbon atoms typically make up (as determined by ASTM D
2887) at least about 10, more typically at least about 15,
even more typically at least about 20, and most typically
at least about 24, weight percent of the white mineral oil.
30 In fact, it is not uncommon for compounds containing 18 or
more carbon atoms to constitute at least about 70, 75, 80,
or even about 85, weight percent of the white mineral oil.

In addition, the white mineral oil comprises up to about 30 weight percent compounds containing 15 or less carbon atoms (as determined by ASTM D 2887). Often the white mineral oil comprises at least about 0.1, more often at least about 0.5, even more often at least about 1, and most often at least about 1.5, weight percent compounds containing 15 or less carbon atoms. Generally, compounds

containing 15 or less carbon atoms account for less than about 27, more commonly less than about 25, and even more typically less than about 23, weight percent of the white mineral oil. In fact, the concentration of compounds containing 15 or less carbon atoms in the white mineral oil can be less than about 15, 10, or even 5 weight percent.

The white mineral oils employed in the present invention preferably have an aromatic content (as determined by ASTM D 2887) of less than about 0.5, more preferably less than about 0.4, even more preferably less than about 0.3, and most preferably less than about 0.2, weight percent.

The pour point of the white mineral oils used in the invention (as determined by ASTM D 97) is generally greater than about -30°C (-22°F), more commonly greater than about -25°C (-13°F), even more commonly greater than about -20°C (-4°F), and most commonly greater than about -15°C (5°F). Usually, the pour point of the white mineral oil is less than about 0°C (32°F).

The flash point of the white mineral oil (as determined by ASTM D 92) is normally greater than about 121.1°C (250°F).

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The viscosity of the white mineral oil at 40°C (104°F) (as measured by ASTM D 445) is ordinarily between about 1 to about 10 centistokes (cst). In one embodiment of the invention, the white mineral oil generally has a viscosity at 40°C (104°F) of about 2 to about 6, and more typically about 3 to about 5, cst. In another embodiment of the invention, the white mineral oil has a viscosity at 40°C (104°F) greater than 6 cst. In this version of the invention, the white mineral oil frequently has a viscosity at 40°C (104°F) of about 6.1 to about 9.5, more often about 6.2 to about 9.25, even more often about 6.3 to about 9, and most often about 6.4 to about 8.75, cst.

The properties of several exemplary white oils within the scope of the present invention are listed in the following Table I.

TABLE I

Exemplary White Oil Properties

	Property	Z	Drakeol 5	Draketex 50	Peneteck°
	Viscosi	ty			
		37.8°C (100°		48/53	38/42
		10°C (104°F)	·	•	
	API 0 15	5.6°C (60°F)	36.9/38.7	37.4/40.2	41.5/43.6
	Specific	Gravity			
	@ 15.6°	C (60°F)'	0.831/0.842	0.824/0.838	0.808/0.818
10	0 25°C	(77°F) ^f		0.817/0.832	
	Flash Po	int ^s , °C/°F	154/310	152/305	129/265
	a.				oil Products
•-					ineral oil meets
15					CFR 172.878
				n food, 178.	
		indirect i	use in food,	and 573.680	for use in
	b.	animal fee	·		
20	ь.				nzoil Products
20				brand white	
		meets the	following FI	DA regulation	ns: 21 CFR
		172.878 co	vering direc	et use in fo	od, 178.3620 for
		indirect u	se in food,	and 573.680	for use in
25	c.	animal fee			
23	.			k of Pennzoi	
		company.	Peneteck bra	and white min	eral oil meets
		rue lollow	ing FDA regu	lations: 21	CFR 172.878
		covering d	irect use in	food, 178.3	620 for
30		indirect u	se in food,	and 573.680	for use in
30	د.	animal fee			
	d.			ce with ASTM	D 445; SUS
			/bolt Univer		
	e.			ce with ASTM	D 445; CST
35	£	denotes cer			
23	f.			ce with ASTM	
	g.	vetermined	in accordance	ce with ASTM	D 92.

TABLE I (continued)

Exemplary White Oil Properties

	Property		Dra	keol 5°	<u>Drak</u>	etex 50	<u>Pene</u>	eteck°	
	Pour Point	h,°C/°F		-9/15		-9/15		-1/30	
5	Initial BE	o',°C/°F	233	9/453	228	.9/444	22	0/428	
	50% BPi, °C	/°F	348	9/660	336	.1/637	289.	4/553	
	Final BPi,	°C/°F	523	9/975	5	15/959	42	5/797	
	Total Para	affins, wt.	&j	35.7		39.3		56.6	
	n-Paraffin	ns, wt. %k ^k		6.32		8.52		24.36	
10	Mononaphth	nenes, wt.	% ^j	42.2		40.5		29.4	
	Polynaphth	nenes, wt.	% ^j	21.8		20.1		13.8	
	Aromatics	, wt. % ^j		0.3		0.1		0.3	
	Carbon Nur	mber Distri	ibuti	loni					
			C#m	Wt.8º	C#m	Wt.%"	<u>C#™</u> W	<u>t.%"</u>	
15			11	0.13	11	0.14	11	0.15	
			12	0.15	12	0.17	12	0.26	
			13	0.26	13	0.35	13	1.29	
			14	0.59	14	0.96	14	5.38	
			15	1.74	15	2.93	15	14.36	
20	•		16	3.97	16	6.15	16	24.21	
			17	7.19	17	9.89	17	30.22	
			18	9.31	18	12.94	18	17.30	
			19	9.73	19	11.83	19	3.70	
,			20	12.60	20	12.05	20	1.00	
25									_
	h.	Determined							
	i.	Determined	l in	accorda	ance	with AS	TM D	2887; BP	
		denotes bo							
	j٠	Determined							
30	k.	Determined	-	_					
	1.	Determined					TM D	2887.	
	m.	C# denotes							
	n.	Wt.% denot	tes v	veight p	perce	nt.			

TABLE I (continued)

Exemplary White Oil Properties

	Propert	EY	<u>Dr</u>	akeol 5°	Dra	ketex 5	D ^b Pen	<u>eteck°</u>
	Carbon	Number	Distribut	ion ^l				
5			<u>C#</u>	Wt.%"	C#m	Wt.80	C#m W	t.%º
			21	14.51	21	12.28	21	0.59
			22	13.68	22	11.04	22	0.42
			23	9.97	23	7.72	23	0.26
			24	5.62	24	4.38	24	0.14
10			25	2.83	25	2.26	25	0.08
			26	1.62	26	1.19	26	0.06
			27	1.21	27	0.73	27	0.06
		•	28	1.00	28	0.50	28	0.05
			29	0.75	29	0.37	>28	0.48
15			30	0.52	30	0.28		
			31	0.38	31	0.23		
			32	0.31	32	0.20		
			33	0.29	33	C.19		
			· 34	0.22	34	0.15		
20			35	0.23	35	0.16		
			36	0.18	36	0.13		
			37	0.16	37	0.12		
			38	0.12	38	0.10	•	
			39	0.13	39	0.10		
25			40	0.09	>39	0.48		
			41	0.08				
			>41	0.43				

TABLE I (continued)

Exemplary White Oil Properties

	Property		Drak	ceol 5	Drake	tex 50	<u>Pene</u>	<u>teck</u>
	n-Paraffin	Carbon	Number	Distr	ibutio	n ^k		
5			<u>C#^j ₩</u>	t. %k	C# Wt	<u>. % C</u>	#j Wt.	<u>ş k</u>
•	•		10	0.01	10	0.14	10	0.01
			11	0.01	11	0.01	11	0
			12	0.02	12	0.02	12	0.21
			13	0.02	13	0.03	13	2.75
10			14	0.06	14	0.14	14	1.99
10			15	0.34	15	0.67	15	1.03
			16	1.00	16	1.73	16	0.33
	•		17	1.60	17	2.41	17	0.11
		•	18	1.19	18	1.54	18	0.04
15			19	1.08	19	1.09	19	0.02
			20	0.50	20	0.42	20	0.01
			21.	0.19	21	0.18		•
			22	0.15	22	0.14		
			23	0.09	23	0.08		
20	•		24	0.03	24	0.03		
			25	0.01	25	0.02		
		٠	26	0.01	26	0.01		
			27	0			•	
			28	0.01				
25			29	0.01				

White mineral oils are generally prepared by removing, by either acid treatment of hydrogenation, unsaturates and aromatics from oil feedstocks. In an exemplary process, the feedstock oil is treated with large quantities of concentrated or fuming sulfuric acid. The reaction product (namely, oil-soluble sulfonic acid), after caustic neutralization and purification, becomes petroleum sulfonate. The unreacted material (i.e., oil feedstock components which do not react with the sulfuric acid) are the saturated aliphatic and naphthenic hydrocarbons which constitute the white mineral oil. Additional details

pertinent to the preparation of white mineral oils are discussed in Meyer, "White Mineral Oil and Petrolatum," Chemical Publishing Co., Inc., Brooklyn, NY (1950) and Biske et al., "White Oil Manufacture," Proceedings Fourth World Petroleum Congress-Section III/A, each of these publications being incorporated herein in their entireties by reference.

Optionally, one or more pour point depressants are employed in the drilling fluid of the present invention 10 to lower the pour point of the white mineral oil. Typical pour point depressants include, but are not limited to, ethylene copolymers, isobutylene polymers, polyaklylnaphthalenes, wax-aromatic condensation products (e.g., wax-naphthalene condensation products, phenol-wax 15 condensation products), polyalkylphenolesters, polyalkylmethacrylates, polymethacrylates, polyalkylated condensed aromatics, alkylaromatic polymers, iminodiimides, and polyalkylstyrene. (The molecular weights for polyaklylnaphthalenes, polyalkylphenolesters, and 20 polyalkylmethacrylates range from about 2,000 to about 10,000.) Because they are non-toxic, ethylene copolymers and isobutylene polymers are the preferred pour point depressants.

Up to about 1 weight percent pour point

depressant is employed. (As used in the specification and claims, the weight percent of the pour point depressant is based upon the weight of the white mineral oil in the drilling fluid, i.e., it is the weight of the pour point depressant divided by the weight of the white mineral oil in the drilling fluid, the quotient being multiplied by 100%.) Preferably, the pour point depressant is employed in a concentration of 0.005 to about 0.5, more preferably about 0.01 to about 0.4, and most preferably about 0.02 to about 0.3, weight percent.

When employed, the pour point depressant is preferably mixed with the white mineral oil and the resulting composition is then combined with any additional additives as described below.

10

One or more emulsifiers, wetting agents, viscosifiers, weighting agents, fluid loss control agents, and shale inhibiting salts are also optionally used in the oil-based drilling fluid of the present invention.

5 Exemplary emulsifiers include, but are not limited to, fatty acids, soaps of fatty acids, and fatty acid derivatives including amido-amines, polyamides, polyamines, esters (such as sorbitan monoleate polyethoxylate, sorbitan dioleate polyethoxylate), imidaxolines, and alcohols.

Typical wetting agents include, but are not limited to, lecithin, fatty acids, crude tall oil, oxidized crude tall oil, organic phosphate esters, modified imidazolines, modified amidoamines, alkyl aromatic sulfates, alkyl aromatic sulfonates, and organic esters of 15 polyhydric alcohols.

Exemplary viscosifiers include, but are not limited to, organophilic clays (e.g., hectorite, bentonite, and attapulgite), oil soluble polymers, polyamide resins, and polycarboxylic acids and soaps. For drilling fluids 20 intended for use in high temperature environments (e.g., where the bottom hole temperature exceeds about 204.4°C (400°F)), it is desirable to employ a sulfonated elastomer polymeric viscosifier. Preferably, the sulfonated elastomer polymeric viscosifier is a neutralized sulfonated 25 elastomer polymer having about 5 to about 100 milliequivalents of sulfonate groups per 100 g of sulfonated polymer. More preferably, the neutralized sulfonated elastomer polymer has about 5 to about 50 milliequivalents, and most preferably about 5 to about 30 30 milliequivalents, of sulfonate groups per 100 g of sulfonated polymer.

Preferably, the sulfonated elastomer polymeric viscosifier is derived from an elastomer polymer selected from the group consisting of ethylene-propylene-diene 35 monomer (EPDM) terpolymers, copolymers of isoprene and styrene sulfonate salt, copolymers of chloroprene and styrene sulfonate salt, copolymers of isoprene and butadiene, copolymers of styrene and styrene sulfonate salt, copolymers of butadiene and styrene sulfonate salt,

copolymers of butadiene and styrene, terpolymers of isoprene, styrene, and styrene sulfonate salt, terpolymers of butadiene, styrene, and styrene sulfonate salt, butyl rubber, partially hydrogenated polyisoprenes, partially hydrogenated natural rubber, partially hydrogenated buna rubber, partially hydrogenated buna rubber, partially hydrogenated polybutadienes, and Neoprene. Methods for obtaining and characteristics of sulfonated elastomer polymers are known to those skilled in the art. See, for example, U.S. Patent 4,447,338, U.S. Patent 4,425,462, U.S. Patent 4,153,588, U.S. Patent 4,007,149, U.S. Patent 3,912,683, and UK Patent Application 2,212,192, these documents being incorporated in their entirety by reference.

Exemplary weighting agents include, but are not limited to barite, iron oxide, gelana, siderite, and calcium carbonate.

Illustrative fluid loss control agents include, but are not limited to, asphaltics (e.g., asphaltenes and 20 sulfonated asphaltenes), amine treated lignite, and gilsonite. For drilling fluids intended for use in high temperature environments (e.g., where the bottom hole temperature exceeds about 204.4°C (400°F)), the fluid loss control agent is preferably a polymeric fluid loss control 25 agent. Exemplary polymeric fluid loss control agents include, but are not limited to, polystyrene, polybutadiene, polyethylene, polypropylene, polybutylene, polyisoprene, natural rubber, butyl rubber, polymers consisting of at least two monomers selected from the group 30 consisting of styrene, butadiene, isoprene, and vinyl carboxylic acid. Individual or mixtures of polymeric fluid loss control agents can be used in the oil base drilling fluid of this invention.

The preferred polymeric fluid loss control agents
are styrene-butadiene copolymers. Characteristics of
exemplary styrene-butadiene copolymers are listed in the
following Table II:

TABLE II

	<u>Characteristic</u>	Exemplary	<u>Styrene-B</u>	<u>utadiene Co</u>	polymers
	Styrene/-				
	Butadiene Ratio	50/50	57/43	90/10	67/33
5	Surfactant Type	Anionic	Anionic	Anionic	Anionic
	Tg, °C	-22	-11	· 76	12
	pН	9.0	6.0	6.5	9.0

All the styrene/butadiene copolymers described in above Table II also contain about 1 to about 3 weight 10 percent copolymerized carboxylic acid (e.g., itaconic acid and acrylic acid).

Common shale inhibiting salts are alkali metal and alkaline-earth metal salts. Calcium chloride and sodium chloride are the preferred shale inhibiting salts.

Typically, the oil-based drilling fluid of the present invention contains the ingredients and properties set forth in the following Table III:

TABLE III

	Ingredient	Typical	<u>Typical</u>
20	Oil, volume %	25 - 85	50-60
	Surfactant (active),		
	pounds per barrel (ppb) bg	1-20	1-10
	Water, volume %	up to 45	10-20
	Weighting agent, ppb	up to 600	150-400
25	Organophilic clay, ppb	0.5-30	1-10
	Non-polymeric fluid loss control		•
	agent, ppb	up to 30	2-15
	Polymeric fluid loss control		
	agent, ppb°	3-12	5-10
30	Sulfonated elastomer polymeric		
	viscosifier, ppbd	0.02-2	0.05-1.5
	Shale inhibiting salt, ppb	up to 60	20-30
	Lime, ppb°	up to 30	1-10
	Property		
35	Density, ppg ^f	7.5-20	9-16
	a. Volume percent is based on the	he total vo	olume of

the drilling fluid.

- As used in the specification and claims, the term b. "surfactant" means a substance that, when present at low concentration in a system, has the property of adsorbing onto the surfaces or 5 interfaces of the system and of altering to a marked degree the surface or interfacial free energies of those surfaces (or interfaces). used in the foregoing definition of surfactant, the term "interface" indicates a boundary between 10 any two immiscible phases and the term "surface" denotes an interface where one phase is a gas. usually air. Exemplary ingredients referred to as surfactants by those skilled in the art include emulsifiers and oil wetting agents. 15 c. The polymeric fluid loss control agent is preferably present in the drilling fluid in a concentration of about 6 to about 9 ppb. d. The sulfonated elastomer polymeric viscosifier is preferably present in the drilling fluid in a concentration of about 0.1 to about 1 ppb. 20 As used in the specification and claims, the term e. "lime" means quicklime (CaO), quicklime precursors, and hydrated quicklime (e.g., slaked lime $(Ca(OH)_2)$. ppg denotes pounds per gallon.
- 25 f.
 - The pounds per barrel (ppb) is based upon the g. final composition of the drilling fluid.

The volumetric ratio of oil to water in the drilling fluid of the present invention can be as low as 30 about 50:50.

Preferably, the weight ratio of the polymeric fluid loss control agent to the sulfonated elastomer polymeric viscosifier is about 1.5:1 to about 50:1, more preferably about 3:1 to about 20:1, and most preferably 35 about 5:1 to about 10:1.

The drilling fluid is preferably prepared by mixing the constituent ingredients in the following order: (a) white mineral oil, (b) organophilic clay, (c)

surfactant, (d) lime, (e) an aqueous solution comprising water and the shale inhibiting salt, (f) non-polymeric fluid loss control agent, (g) weighting agent, (h) polymeric fluid loss control agent, and (i) sulfonated elastomer polymeric viscosifier.

While it is preferred that the entire base oil of the drilling fluid consist solely of the white mineral oil (and any optional pour point depressant), the white mineral oil can constitute a portion of the base oil. Preferably, 10 the white mineral oil constitutes at least about 50, more preferably at least about 75, even more preferably at least about 90, and most preferably at least about 95, weight percent of the base oil. (As used in the specification and claims, the term "weight percent" in the context of the 15 concentration of the white mineral oil in the base oil means the weight of the white oil used in the drilling fluid divided by the weight of all oils employed to formulate the base oil component of the drilling fluid, the quotient being multiplied by 100 percent.) It follows that 20 the base oil can consist of about 96, 97, 98, 99, or more, weight percent white mineral oil.

EXAMPLES

The following examples (which are intended to illustrate and not limit the invention, the invention being defined by the claims) demonstrate the preparation and properties of exemplary drilling fluids within the scope of the present invention.

EXAMPLES 1-3

30

Preparation And Initial Properties Of Drilling Fluids

Low toxicity, oil-based drilling fluids were
prepared using non-toxic, food grade white oils as the
base.

Each drilling fluid was an invert emulsion prepared by (a) initially agitating about 240 ml of a nontoxic, food grade white oil for about 1 minute using a
Multi-Mixer brand blender and (b) then sequentially adding the following ingredients (with continuous mixing for about one minute after the addition of each material): (i) about 6 g of Invermul NT brand primary emulsifier (available from

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Baroid Drilling Fluids, Inc. (Baroid)); (ii) about 8 g of lime (calcium hydroxide); and (iii) about 4 g of Duratone HT brand fluid-loss preventing agent (i.e., a chemically modified lignite (humate) available from Baroid).

Subsequently, about 39 ml of fresh water was added to the above mixture and the resulting composition was mixed for about ten minutes. Then, about 11 g of Geltone II brand amine-treated bentonite (available from Baroid) was added and the resulting matter was agitated for about 15 minutes.

Thereafter, the following materials were added in sequence, with about 5 minutes of mixing after the addition of each of the materials: (i) about 2 g of EZ Mul NT brand secondary emulsifier (available Baroid); (ii) about 210 g of powdered barite (a non-toxic weighting agent); (iii) about 24 g of calcium chloride dihydrate (to provide salinity to the water phase without water wetting the barite); and (iv) about 20 g of a powdered clay (composed of about 35 weight percent smectite and about 65 weight percent kaolinite) to simulate drilled formation particles.

The rheological properties of each of the three initial drilling fluids was determined at about 65.6°C (150°F) and are set forth below in Table A.

35

TABLE A

Initial Rheological Properties

			Drillin	g Fluid
	Parameter	A*	B ^b	C°
5	Fann Viscometer		•	
	Dial Reading at:			
	600 RPM	100	93	60
	300 RPM	62	55	35
-	3 RPM	12	11	6
10	Plastic Viscosity,			
	N-sec/m²/centipoise	0.038/38	0.038/38	0.025/25
	Yield Point',			
	$kg/9.29 m^2/lb/100 ft^2$	10.9/24	7.7/17	4.5/10
	10 Second Gel Strength,	,		
15	$kg/9.29 m^2/lb/100 ft^2$	6.8/15	5.9/13	3.2/7
	10 Minute Gel Strength,			
	$kg/9.29 m^2/lb/100 ft^2$	10.0/22	9.1/20	5.0/11
	Density,			
	<pre>lb/gal/kg/l lb./gallon</pre>	1.4/11.8	1.4/11.8	1.4/11.8
20	HTHP Filtrate at 148.9°C			
	(300°F) and 3548.7			
	kpascal (500 psig) ^b , ml	9	10	11
	Emulsion Stability ⁱ , volt	1203	1258	987
			 :	
25	a. "A" is Drakeol			•
	b. "B" is Draketex			
	c. "C" is Peneteck			
	d. Dial readings w		_	
	motor driven vi		_	
30	Practice - Stan			
	Drilling Fluids	- •		
	(RP13B), Twelft	•	•	•
	American Petrol (hereinafter re			
	(Heternatuer re	retten to as	WET.,)'	vayes /-3,

sections 2.4 to 2.5.

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- e. PV was determined in accordance with the procedure and calculations discussed in API, page 9, sections 2.5 to 2.6.
- f. YP was determined in accordance with the procedure and calculations discussed in API, page 9, sections 2.5 to 2.6.
 - g. Gel strength for 10 seconds and 10 minutes was determined in accordance with the procedure discussed in API, page 9, section 2.5, paragraphs f and g, respectively.
 - h. HTHP (High-Temperature/High-Pressure) Filtrate was determined in accordance with the procedure discussed in API, page 12, section 3.5.
- i. Emulsion stability was measured according to the
 procedure described in API, page 28.

EXAMPLE 4-6

Age Tested Drilling Fluid Properties Aging Methodology

Each of the three above-prepared drilling fluids
was age tested in accordance with the following protocol.
A portion of the drilling fluid was transferred to a
stainless steel vessel which was then pressurized with
nitrogen to about 790.8 kpascal (100 psig). The
pressurized, drilling fluid-containing vessel was placed on
a rolling apparatus inside an oven set at about 65.6°C
(150°F) for about 16 hours of aging. The rheological
properties for each of the aged drilling fluids are listed
in the following Table B.

TABLE B
Aged Rheological Properties

		D;	rilling Flu	<u>id</u>
	<u>Parameter</u>	A*	B _p	C°
5	Fann Viscometer			
	Dial Reading at:		•	
	600 RPM	110	100	68
	300 RPM	66	60	39
	3 RPM	12	50	6
10	Plastic Viscosity,			
	N-sec/m²/centipoise	0.044/44	0.040/40	0.029/29
	Yield Point ^f ,			
	$kg/9.29 m^2/lb/100 ft^2$	10.0/22	9.1/20	4.5/10
	10 Second Gel Strengths,			
15	$kg/9.29 m^2/lb/100 ft^2$	6.8/15	6.4/14	3.6/8
	10 Minute Gel Strength,			
	$kg/9.29 m^2/lb/100 ft^2$	10.9/24	10.4/23	6.4/14
	Density,			
	lb/gal/kg/l lb./gallon	1.4/11.8	1.4/11.8	1.4/11.8
20	Filtrate at 148.9°C (300°)	F)		
	and 3548.7 kpascal			
	(500 psig) ^b , ml	12	7	16
	Emulsion Stability, volt	1454	1502	1227

25 a-i. See above Table A.

The initial and aged rheological properties of each of the three exemplary oil-based drilling fluids of the present invention are appropriate for an oil-based drilling fluid. For example, the low initial and aged plastic viscosities indicate that each of the exemplary drilling muds can maintain low pumping pressures and the high initial and aged yield points and gel strengths show that each drilling mud can effectively suspend solids to clean boreholes.

Although the present invention has been described in detail with reference to some preferred versions, other versions are possible. For example, the white mineral oil

can also be employed as the base oil component in other wellbore fluids. (As used in the specification and claims, the term "wellbore fluid" means a fluid used while conducting pay zone drilling, underreaming, drilling in,

5 plugging back, sand control, perforating, gravel packing, chemical treatment, hydraulic fracturing, cleanout, well killing, tubing and hardware replacement, and zone selective operations (e.g., well completion operations) as well as a fluid employed as a packer fluid.) In addition to the base oil, the wellbore fluids contain one or more additional ingredients such as proppants suitable for use in hydraulically fracturing subterranean formations, particulate agents suitable for use in forming a gravel pack, viscosifiers, organophilic clays, and fluid loss control agents.

Common proppants suitable for use in hydraulic fracturing procedures are quartz sand grains, tempered glass beads, sintered bauxite, resin coated sand, aluminum pellets, and nylon pellets. Generally, the proppants are employed in the wellbore fluids intended for use as hydraulic fracturing fluids and are used in concentrations of roughly about 1 to about 10 pounds per gallon of the wellbore fluid. The proppant size is typically smaller than about 2 mesh on the U.S. Sieve Series scale, with the exact size selected being dependent on the particular type of formation to be fractured, the available pressure and pumping rates, as well as other factors known to those skilled in the art.

Typical particulate agents employed in the

wellbore fluids used as gravel packing fluids include, but
are not limited to, quartz sand grains, glass beads,
synthetic resins, resin coated sand, walnut shells, and
nylon pellets. The gravel pack particulate agents are
generally used in concentrations of about 1 to about 20

pounds per gallon of the wellbore fluid. The size of the
particulate agent employed depends on the type of
subterranean formation, the average size of formation
particles, and other parameters known to those skilled in

the art. Generally, particulate agents of about 8 to about 70 mesh on the U.S. Sieve Series scale are used.

Illustrative viscosifiers, organophilic clays, and fluid loss control agents optionally used in wellbore 5 fluids and their concentrations are the same as discussed above in connection with drilling fluids.

The wellbore fluids are prepared by combining the white mineral oil and pour point depressant (when used) and adding any additional additive (e.g., hydraulic fracturing 10 proppants, gravel pack particulate agents, viscosifiers, fluid loss control agents, and organophilic clays) is usually added to resulting combination.

The specific techniques used when employing the wellbore fluid are determined by its intended use and are 15 analogous to methodologies employed when using prior art wellbore fluids for corresponding completion or work-over operations. For example, when the wellbore fluid is employed as a gravel packing fluid, it is typically injected into the formation in accordance with the 20 procedure discussed in U.S. Patent 4,552,215, this patent being incorporated herein in its entirety by reference.

When employed as a fracturing fluid, the wellbore fluid of the present invention is usually injected into the formation using procedures analogous to those disclosed in 25 U.S. Patent 4,488,975, U.S. Patent 4,553,601, Howard et al., Hydraulic Fracturing, Society of Petroleum Engineers of the American Institute of Mining, Metallurgical, and Petroleum Engineers, Inc., New York, NY (1970), and Allen et al., Production Operations, Well completions, Workover, 30 and Stimulation, 3rd Edition, volume 2, Oil & Gas Consultants International, Inc., Tulsa, Oklahoma (1989) (Allen), chapter 8, these publications being incorporated

When employed in a perforating operation, the 35 wellbore fluid of the present invention is used according to the methodologies disclosed in chapter 7 of Allen, referenced above.

herein in their entirety by reference.

Techniques for using packer fluids and well killing fluids, such as those discussed in chapter 8 of Allen, are also applicable to the wellbore fluid of the present invention.

In view of the foregoing numerous other embodiments, the spirit and scope of the appended claims should not necessarily be limited to the description of the preferred versions contained herein.

Attachment A

Normal Hydrocarbons In Petroleum Products

By Gas Chromatography

Normal hydrocarbons can be determined in petroleum products using gas chromatography with a flame ionization detector (FID). A capillary column 12.5 meters long, 0.32 mm id., and with a 0.5 μ m film of Ohio Valley brand OV-1 silicon polymer or an equivalent stationary phase is used.

10 Column conditions are set as follows:

Injector: 350°C

Split flow set at 170 ml/min.
Head pressure set at 170.27 kpascal (10

psig)

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15 Detector: FID at 350°C

Oven: 100 degrees 2 minutes then

8 degrees/min. to 300 degrees.

Hold at 300 degrees for 15 minutes.

Injection size is 1 μ L.

20 Sample preparation:

Internal standard: 0.2xxx g p-terphenyl is dissolved in

100 mL toluene. This does not dissolve quickly and may take

overnight.

25 Sample: A 1 dram vial is tarred and 0.5xxx g

of sample is weighed into the vial.

2.00 mL of the internal standard
solution is added to this. Shake
well to completely dissolve the

sample. Warming may be required for

some samples.

Attachment A (continued)

Analysis:

Inject 1 μ L into the gas chromatograph and start the run. Check that all normal hydrocarbons and p-terphenyl are correctly identified. The relative response factor between p-terphenyl and the normal hydrocarbons should have been determined in a previous run from a sample prepared to have a known composition. Integrate all identified peaks from valley to valley. The amount of each hydrocarbon is calculated from the relative peak areas as compared to the internal standard and the amount of the internal standard.

CLAIMS

- 1. A wellbore fluid comprising:
- (a) at least one additive selected from the group consisting of emulsifiers, wetting agents, viscosifiers,
 5 weighting agents, fluid loss control agents, proppants for use in hydraulically fracturing subterranean formations, and particulate agents for use in forming a gravel pack;
 and
- (b) a white mineral oil having (i) an API gravity 10 at 15.6°C (60°F) greater than 35, (ii) a content of compounds containing 14 or more carbon atoms of at least about 95 weight percent, and (iii) a pour point of at least about -30°C (-22°F).
- The wellbore fluid of claim 1 wherein the white
 mineral oil has a n-paraffinic content greater than 5
 weight percent.
 - 3. The wellbore fluid of claims 1 or 2 wherein the white mineral comprises at least about 10 weight percent compounds containing 18 or more carbon atoms.
- 20 4. The wellbore fluid of claims 1 or 2 wherein the white mineral comprises at least about 15 weight percent compounds containing 18 or more carbon atoms.
- 5. The wellbore fluid of claims 1 or 2 wherein the white mineral comprises at least about 20 weight percent compounds containing 18 or more carbon atoms.
 - 6. The wellbore fluid of claims 1 or 2 wherein the white mineral comprises at least about 96 weight percent compounds containing 14 or more carbon atoms.
- 7. The wellbore fluid of claims 1 or 2 wherein the 30 white mineral comprises at least about 97 weight percent compounds containing 14 or more carbon atoms.
 - 8. The wellbore fluid of claims 1 or 2 wherein the white mineral comprises at least about 98 weight percent compounds containing 14 or more carbon atoms.
- 35 9. The wellbore fluid of claims 1 to 8 wherein the white mineral oil has a characterization factor greater than 12.

10. The wellbore fluid of claims 1 to 8 wherein the white mineral oil has a characterization factor of at least about 12.1.

- 11. The wellbore fluid of claims 1 to 8 wherein the
- 5 white mineral oil has a characterization factor of at least about 12.2.
 - 12. The wellbore fluid of claims 1 to 11 wherein the white mineral oil has an API gravity at 15.6°C (60°F) greater than about 35.1.
- 10 13. The wellbore fluid of claims 1 to 11 wherein the white mineral oil has an API gravity at 15.6°C (60°F) greater than about 35.25.
 - 14. The wellbore fluid of claims 1 to 11 wherein the white mineral oil has an API gravity at 15.6°C (60°F)
- 15 greater than about 35.5.
 - 15. The wellbore fluid of claims 1 to 14 further comprising a pour point depressant.
 - 16. A drilling system comprising:
 - (A) at least one subterranean formation;
- 20 (B) a borehole penetrating a portion of at least one of the subterranean formations;
 - (C) a drill bit suspended in the borehole; and
 - (D) a drilling fluid located in the borehole and proximate the drill bit,
- wherein the drilling fluid is the wellbore fluid of claim 1 and at least one additive is selected from the group consisting of emulsifiers, wetting agents, viscosifiers, weighting agents, and fluid loss control agents.
 - 17. A method for drilling a borehole in a
- 30 subterranean formation, the method comprising the steps of:
 - (A) rotating a drill bit at the bottom of the borehole; and
 - (B) introducing a drilling fluid into the borehole (α) to pick up drill cuttings and (β) to carry at
- 35 least a portion of the drilling cuttings out of the borehole,
 - wherein the drilling fluid is the wellbore fluid of claim 1 and at least one additive is selected from the group

consisting of emulsifiers, wetting agents, viscosifiers, weighting agents, and fluid loss control agents.

- 18. A method for the drilling or completion or workover of a well, the method being characterized by the step

 5 of using a white mineral oil having (i) an API gravity at
 15.6°C (60°F) greater than 35, (ii) a content of compounds
 containing 14 or more carbon atoms of at least about 95
 weight percent, and (iii) a pour point of at least about 30°C (-22°F).
- 10 19. A natural resource system comprising:
 - (a) a subterranean formation;
 - (b) a well penetrating at least a portion of the subterranean formation; and
- (c) the wellbore fluid of claim 1 present in at 15 least a portion of the well.
 - 20. A natural resource system comprising:
 - (a) a subterranean formation;
 - (b) a wellbore penetrating at least a portion of the subterranean formation;
- 20 (c) a casing positioned within at least a portion of the wellbore; and
 - (d) the wellbore fluid of claim 1 present in at least a portion of the area between the surface of the wellbore and the outside surface of the casing.
- 25 21. A natural resource system comprising:
 - (a) a subterranean formation;
 - (b) a wellbore penetrating at least a portion of the subterranean formation;
- (c) a casing positioned within at least a portion 30 of the wellbore; and
 - (d) the wellbore fluid of claim 1 present in at least a portion of the subterranean formation proximate the wellbore.

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A. CLASSII IPC 5	FICATION OF SUBJECT MATTER C09K7/06			
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